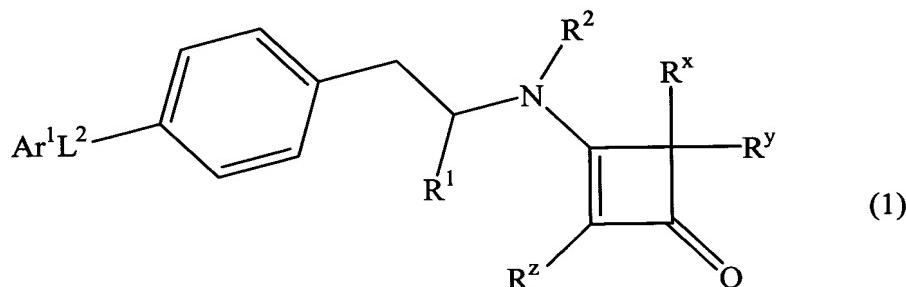


This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (currently amended) A process for the preparation of a compound of formula (1):



wherein:

Ar^1 is an optionally substituted aromatic or heteroaromatic group;

L^2 is a linker group selected from $-\text{N}(\text{R}^4)-$ [where R^4 is a hydrogen atom or an optionally substituted straight or branched C₁₋₆alkyl group], $-\text{CON}(\text{R}^4)-$ and $-\text{S}(\text{O})_2\text{N}(\text{R}^4)-$;

R^4 is a hydrogen atom or an optionally substituted straight or branched C₁₋₆alkyl group;

R^1 is a carboxylic acid ($-\text{CO}_2\text{H}$) or a derivative or biostere thereof an acyclic or cyclic carboxylic acid ester, an amide, tetrazole, phosphonic acid, phosphinic acid, sulphonic acid, sulphinic acid, boronic acid, or an acylsulphonamide group;

R^2 is a hydrogen atom or a C₁₋₆alkyl group;

R^x , R^y and R^z , which may be the same or different, are each an atom or group $-\text{L}^1(\text{Alk}^1)_n(\text{R}^3)_v$ in which , or R^z is $-\text{L}^1(\text{Alk}^1)_n(\text{R}^3)_v$ and R^x and R^y are joined together to form an optionally substituted spiro linked cycloaliphatic or heterocycloaliphatic group;

L^1 is a covalent bond or a linker atom or group an $-\text{O}-$, $-\text{S}-$, or $-\text{Se}-$ atom or an $-\text{C}(\text{O})-$, $-\text{C}(\text{O})\text{O}-$, $-\text{OC}(\text{O})-$, $-\text{C}(\text{S})-$, $-\text{S}(\text{O})-$, $-\text{S}(\text{O})_2-$, $-\text{N}(\text{R}^8)-$, $-\text{CON}(\text{R}^8)-$, $-\text{OC}(\text{O})\text{N}(\text{R}^8)-$, $-\text{CSN}(\text{R}^8)-$, $-\text{N}(\text{R}^8)\text{CO}-$, $-\text{N}(\text{R}^8)\text{C}(\text{O})\text{O}-$, $-\text{N}(\text{R}^8)\text{CS}-$, $-\text{S}(\text{O})_2\text{N}(\text{R}^8)-$, $-\text{N}(\text{R}^8)\text{S}(\text{O})_2-$, $-\text{N}(\text{R}^8)\text{O}-$, $-\text{ON}(\text{R}^8)-$, $-\text{N}(\text{R}^8)\text{N}(\text{R}^8)-$, $-\text{N}(\text{R}^8)\text{CON}(\text{R}^8)-$, $-\text{N}(\text{R}^8)\text{CSN}(\text{R}^8)-$, or $-\text{N}(\text{R}^8)\text{SO}_2\text{N}(\text{R}^8)-$ group;

R⁸ is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group;

Alk¹ is an optionally substituted aliphatic chain or an optionally substituted heteroaliphatic chain containing one to four -O- or -S- atoms or -C(O)-, -C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸), -N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-, -N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- groups that interrupt or are at the terminus of the aliphatic chain;

R³ is a hydrogen or halogen atom or group selected from -OR^{3a} [where R^{3a} is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group or C₃-8cycloalkyl group], -SR^{3a}, -CN and an optionally substituted cycloaliphatic, heterocycloaliphatic, polycycloaliphatic, heteropolycycloaliphatic, aromatic or heteroaromatic group; ;

R^{3a} is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group or C₃-8cycloalkyl group;

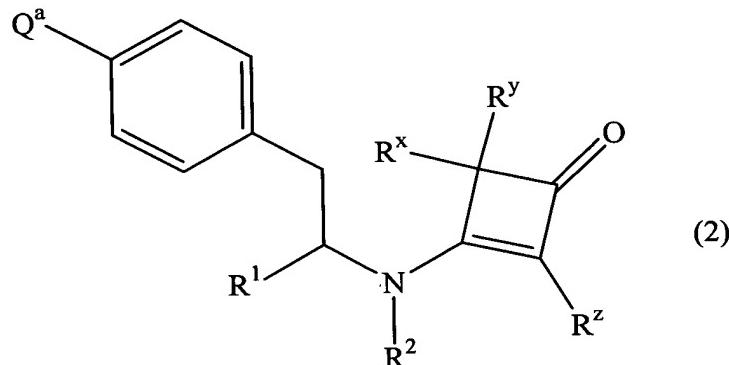
n is zero or the integer 1; and

v is the integer 1, 2 or 3;

provided that when n is zero and L¹ is a covalent bond, v is the integer 1;

or R^Z is an atom or group as previously defined and R^X and R^Y are joined together to form an optionally substituted spiro linked cycloaliphatic or heterocycloaliphatic group; and the salts, solvates, hydrates and N-oxides thereof;

which comprises reacting a compound of formula (2):



wherein:

Q^a is a group $-N(R^4)H$;

and the salts, solvates, hydrates and N-oxides thereof;

with a compound Ar^1W wherein

W is a group selected from X^1 (~~wherein X^1 is a leaving atom or group~~), $-COX^2$

(~~wherein X^2 is a halogen atom or a -OH group~~) and

$-SO_2X^3$ (~~in which X^3 is a halogen atom~~);

X^1 is a leaving atom or group;

X^2 is a halogen atom or a -OH group; and

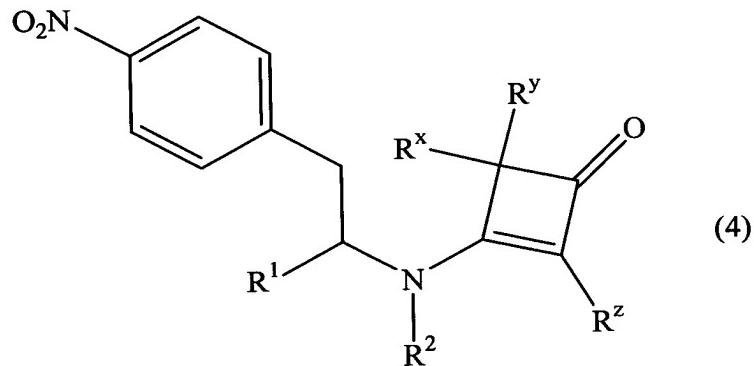
X^3 is a halogen atom.

2. (original) A process according to Claim 1 wherein the reaction is carried out in a solvent in the presence of an acid when W is the group X^1 .

3. (previously presented) A process according to Claim 2 wherein the solvent is selected from an alcohol, ether, acetic acid, water, acetonitrile, substituted amide and ester.

4. (original) A process according to Claim 2 wherein the reaction is carried out in an alcohol in the presence of an acid catalyst.

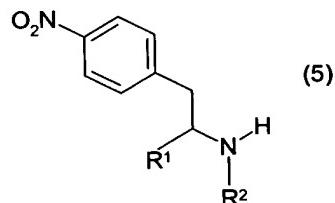
5. (original) A process according to Claim 1 wherein the reaction is carried out in the presence of a base, an organic amine or a cyclic amine and an organic solvent when W is the group COX^2 and X^2 is a halogen atom.
6. (previously presented) A process according to Claim 5 wherein the organic solvent is selected from a halogenated hydrocarbon, a dipolar aprotic solvent, an ether and an ester.
7. (original) A process according to Claim 1 wherein the reaction is carried out in the presence of a condensing agent and a halogenated hydrocarbon, dipolar aprotic or an ether solvent when W is the group CO_2H .
8. (original) A process according to Claim 1 wherein the reaction is carried out in the presence of a base, an organic amine or a cyclic amine and a halogenated hydrocarbon, dipolar aprotic or an ether solvent when W is the group SO_2X^3 .
9. (previously presented) A process according to claim 1 wherein the compound of formula (2) is prepared by reduction of a compound of formula (4):



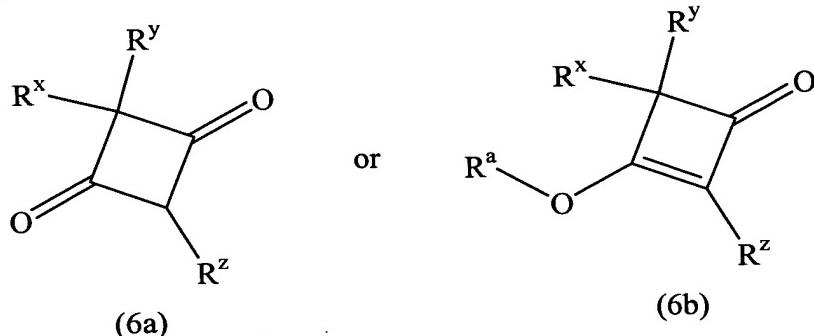
10. (original) A process according to Claim 9 wherein the reduction is carried out by catalytic hydrogenation or by chemical reduction.

11. (previously presented) A process according to Claim 1 wherein R⁴ is a hydrogen atom.

12. (original) A process according to Claim 9 wherein the compound of formula (4) is prepared by reaction of a compound of formula (5):



with a compound of formula (6a) or (6b):



wherein R^a represents a C₁-6alkyl group or a silyl group.

13. (original) A process according to Claim 12 wherein the reaction is carried out in the presence of an organic solvent.

14. (previously presented) A process according to Claim 13 wherein the solvent is selected from an aromatic hydrocarbon, a halogenated hydrocarbon and an ester.

15. (currently amended) A process according to Claim 1 wherein R¹ is the group -CO₂Alk⁷; and

Alk⁷ is a straight or branched optionally substituted C₁-8alkyl group, an optionally substituted C₂-8alkenyl group, an optionally substituted C₂-8alkynyl group, an optionally substituted C₃-8cycloalkyl group, an optionally substituted C₃-8heterocycloalkyl group, an

optionally substituted C₃-8cycloalkylC₁-8alkyl group, an optionally substituted C₃-8heterocycloalkylC₁-8alkyl group, an optionally substituted C₁-6alkyloxyC₁-6alkyl group, an optionally substituted hydroxyC₁-6alkyl group, an optionally substituted C₁-6alkylthioC₁-6alkyl group, an optionally substituted C₁-6alkylsulfinylC₁-6alkyl group, an optionally substituted C₃-8cycloalkyloxyC₁-6alkyl group, an optionally substituted C₃-8cycloalkylthioC₁-6alkyl group, an optionally substituted C₃-8cycloalkylsulfinylC₁-6alkyl group, an optionally substituted C₃-8cycloalkylsulfonylC₁-6alkyl group, an optionally substituted C₁-6alkyloxycarbonylC₁-6alkyl group, an optionally substituted C₁-6alkyloxycarbonyloxyC₁-6alkyl group, an optionally substituted C₁-6alkyloxycarbonyloxyC₁-6alkenyl group, an optionally substituted C₃-8cycloalkyloxycarbonyloxyC₁-6alkyl group, an optionally substituted N-di-C₁-8alkylaminoC₁-8alkyl group, an optionally substituted N-C₆-12aryl-N-C₁-6alkylaminoC₁-6alkyl group, an optionally substituted N-di-C₁-8alkyl-carbamoylC₁-galkyl group, an optionally substituted C₆-12arylC₁-6alkyl group, an optionally substituted heteroC₆-10arylC₁-6alkyl group, an optionally substituted C₆-12aryl group, an optionally substituted C₆-12aryloxyC₁-galkyl group, an optionally substituted C₆-12arylthioC₁-8alkyl group, an optionally substituted C₆-12arylsulfinylC₁-8alkyl group, an optionally substituted C₆-12arylsulfonylC₁-8alkyl group, an optionally substituted C₁-8alkanoyloxyC₁-8alkyl group, an optionally substituted C₄-8imidoC₁-8alkyl group, an optionally substituted C₆-12aroyloxyC₁-8alkyl group, or a triglyceride.

16. (canceled)

17. (currently amended) A process according to Claim 16 Claim 1 which comprises hydrolysing a compound of formula (1) in which R¹ is -CO₂Alk⁷ and Alk⁷ is a straight or branched optionally substituted C₁-8alkyl group, an optionally substituted C₂-8alkenyl group, an optionally substituted C₂-8alkynyl group, an optionally substituted C₃-8cycloalkyl

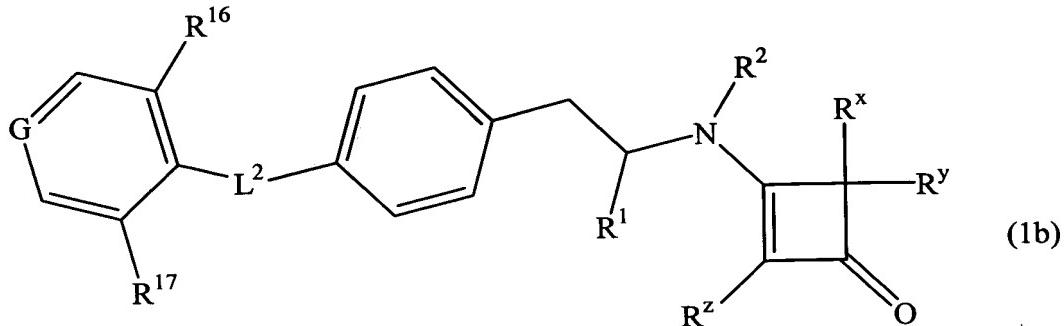
group, an optionally substituted C₃-heterocycloalkyl group, an optionally substituted C₃-cycloalkylC₁-8alkyl group, an optionally substituted C₃-heterocycloalkylC₁-8alkyl group, an optionally substituted C₁-6alkyloxyC₁-6alkyl group, an optionally substituted hydroxyC₁-6alkyl group, an optionally substituted C₁-6alkylthioC₁-6alkyl group, an optionally substituted C₁-6alkylsulfinylC₁-6alkyl group, an optionally substituted C₁-6alkylsulfonylC₁-6alkyl group, an optionally substituted C₃-8cycloalkyloxyC₁-6alkyl group, an optionally substituted C₃-8cycloalkylthioC₁-6alkyl group, an optionally substituted C₃-8cycloalkylsulfinylC₁-6alkyl group, an optionally substituted C₃-8cycloalkylsulfonylC₁-6alkyl group, an optionally substituted C₁-6alkyloxycarbonylC₁-6alkyl group, an optionally substituted C₁-6alkyloxycarbonylC₁-6alkenyl group, an optionally substituted C₁-6alkyloxycarbonyloxyC₁-6alkenyl group, an optionally substituted C₃-8cycloalkyloxycarbonyloxyC₁-6alkyl group, an optionally substituted N-di-C₁-8alkylaminoC₁-8alkyl group, an optionally substituted N-C₆-12aryl-N-C₁-6alkylaminoC₁-6alkyl group, an optionally substituted N-di-C₁-8alkyl-carbamoylC₁-galkyl group, an optionally substituted C₆-12arylC₁-6alkyl group, an optionally substituted heteroC₆-10arylC₁-6alkyl group, an optionally substituted C₆-12aryl group, an optionally substituted C₆-12aryloxyC₁-8alkyl group, an optionally substituted C₆-12arylthioC₁-8alkyl group, an optionally substituted C₆-12aryl sulfinylC₁-8alkyl group, an optionally substituted C₆-12arylsulfonylC₁-8alkyl group, an optionally substituted C₁-8alkanoyloxyC₁-8alkyl group, an optionally substituted C₄-8imidoC₁-8alkyl group, an optionally substituted C₆-12aroyloxyC₁-8alkyl group, or a triglyceride,

to produce a compound of formula (1) in which R¹ is -CO₂H.

18. (currently amended) A process according to Claim 16 Claim 1 which comprises esterifying a compound of formula (1) in which R¹ is -CO₂H to produce a compound of formula (1) in which R¹ is -CO₂Alk⁷ and Alk⁷ is a straight or branched optionally substituted C₁-8alkyl group, an optionally substituted C₂-8alkenyl group, an optionally substituted C₂-

galkynyl group, an optionally substituted C₃-cycloalkyl group, an optionally substituted C₃-heterocycloalkyl group, an optionally substituted C₃-cycloalkylC₁-galkyl group, an optionally substituted C₃-heterocycloalkylC₁-galkyl group, an optionally substituted C₁-6alkyloxyC₁-6alkyl group, an optionally substituted hydroxyC₁-6alkyl group, an optionally substituted C₁-6alkylthioC₁-6alkyl group, an optionally substituted C₁-6alkylsulfinylC₁-6alkyl group, an optionally substituted C₃-8cycloalkyloxyC₁-6alkyl group, an optionally substituted C₃-8cycloalkylthioC₁-6alkyl group, an optionally substituted C₃-8cycloalkylsulfinylC₁-6alkyl group, an optionally substituted C₃-8cycloalkylsulfonylC₁-6alkyl group, an optionally substituted C₁-6alkyloxycarbonylC₁-6alkyl group, an optionally substituted C₁-6alkyloxycarbonylC₁-6alkenyl group, an optionally substituted C₁-6alkyloxycarbonyloxyC₁-6alkenyl group, an optionally substituted C₃-8cycloalkyloxycarbonyloxyC₁-6alkyl group, an optionally substituted N-di-C₁-8alkylaminoC₁-8alkyl group, an optionally substituted N-C₆-12aryl-N-C₁-6alkylaminoC₁-6alkyl group, an optionally substituted N-di-C₁-8alkyl-carbamoylC₁-8alkyl group, an optionally substituted C₆-12arylC₁-6alkyl group, an optionally substituted heteroC₆-10arylC₁-6alkyl group, an optionally substituted C₆-12aryl group, an optionally substituted C₆-12aryloxyC₁-8alkyl group, an optionally substituted C₆-12arylthioC₁-8alkyl group, an optionally substituted C₆-12arylsulfinylC₁-8alkyl group, an optionally substituted C₆-12arylsulfonylC₁-8alkyl group, an optionally substituted C₁-8alkanoyloxyC₁-8alkyl group, an optionally substituted C₄-8imidoC₁-8alkyl group, an optionally substituted C₆-12aroyloxyC₁-8alkyl group, or a triglyceride.

19. (currently amended) A process according to Claim 1 for the preparation of compounds of formula (1b):



wherein

-G= is -CR¹⁸=, -N= or -N(O)=;

R¹⁶, R¹⁷ and R¹⁸, which may be the same or different, are each a hydrogen atom or an atom or group -L³(Alk²)_tL⁴(R⁵)_u;

L³ and L⁴ are, independently, a covalent bond, an -O- or -S- atom, or a -C(O)-, -C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸)-, -N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-, -N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- group;

R⁸ is a hydrogen atom or an optionally substituted straight or branched C₁₋₆alkyl group;

t is zero or the integer 1;

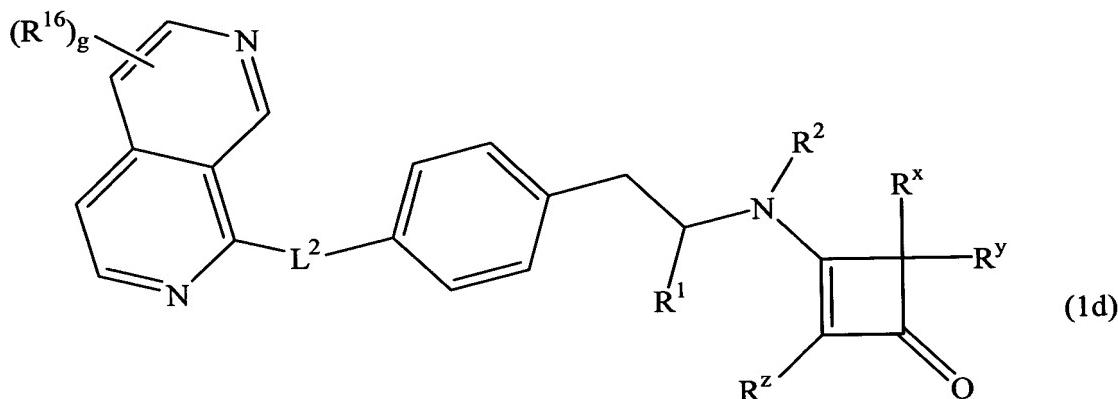
u is an integer 1, 2 or 3;

Alk² is an optionally substituted aliphatic chain or an optionally substituted heteroaliphatic chain containing one to four -O- or -S- atoms or -C(O)-, -C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸)-, -N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-, -N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- groups that interrupt or are at the terminus of the aliphatic chain;

R⁵ is a hydrogen or halogen atom or an optionally substituted C₁₋₆alkyl, optionally substituted C₃₋₈cycloalkyl, -OR⁶, -SR⁶, -NR⁶R⁷, -NO₂, -CN, -CO₂R⁶, -SO₃H, -SOR⁶, -SO₂R⁶, -SO₃R⁶, -OCO₂R⁶, -CONR⁶R⁷, -OCONR⁶R⁷, -CSNR⁶R⁷, -COR⁶, -OCOR⁶,

-N(R⁶)COR⁷, -N(R⁶)CSR⁷, -SO₂N(R⁶)(R⁷), -N(R⁶)SO₂R⁷, N(R⁶)CON(R⁷)(R¹⁹),
-N(R⁶)CSN(R⁷)(R¹⁹), or -N(R⁶)SO₂N(R⁷)(R¹⁹) group; and
R⁶, R⁷, and R¹⁹ are, independently, a hydrogen atom or an optionally substituted C₁-
6alkyl or C₃-8cycloalkyl group;
provided that when t is zero and each of L³ and L⁴ is a covalent bond, then u is the
integer 1 and R⁵ is other than a hydrogen atom;
and the salts, solvates, hydrates and N-oxides thereof.

20. (currently amended) A process according to Claim 1 for the preparation of compounds of formula (1d):



wherein

g is the integer 1, 2, 3 or 4;

R¹⁶, is an atom or group -L³(Alk²)_tL⁴(R⁵)_u;

L³ and L⁴ are, independently, a covalent bond, an -O- or -S- atom, or a -C(O)-,
-C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸)-,
-N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-,
-N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- group;

R⁸ is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group;

t is zero or the integer 1;

u is an integer 1, 2 or 3;

Alk² is an optionally substituted aliphatic chain or an optionally substituted heteroaliphatic chain containing one to four -O- or -S- atoms or -C(O)-, -C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸)-, -N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-, -N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- groups that interrupt or are at the terminus of the aliphatic chain;

R⁵ is a hydrogen or halogen atom or an optionally substituted C₁-6alkyl, optionally substituted C₃-8cycloalkyl, -OR⁶, -SR⁶, -NR⁶R⁷, -NO₂, -CN, -CO₂R⁶, -SO₃H, -SOR⁶, -SO₂R⁶, -SO₃R⁶, -OCO₂R⁶, -CONR⁶R⁷, -OCONR⁶R⁷, -CSNR⁶R⁷, -COR⁶, -OCOR⁶, -N(R⁶)COR⁷, -N(R⁶)CSR⁷, -SO₂N(R⁶)(R⁷), -N(R⁶)SO₂R⁷, N(R⁶)CON(R⁷)(R¹⁹), -N(R⁶)CSN(R⁷)(R¹⁹), or -N(R⁶)SO₂N(R⁷)(R¹⁹) group; and

R⁶, R⁷, and R¹⁹ are, independently, a hydrogen atom or an optionally substituted C₁-6alkyl or C₃-8cycloalkyl group;

provided that when t is zero and each of L³ and L⁴ is a covalent bond, then u is the integer 1 and R⁵ is other than a hydrogen atom;
and the salts, solvates, hydrates and N-oxides thereof.

21. (previously presented) A process according to Claim 1 for the preparation of:

ethyl (2S)-2-[(2-bromo-3-oxospiro[3.5]non-1-en-1-yl)amino]-3-{4-[(3,5-dichloroisonicotinoyl)amino]phenyl}propanoate;
and the salts, solvates, hydrates and N-oxides thereof.

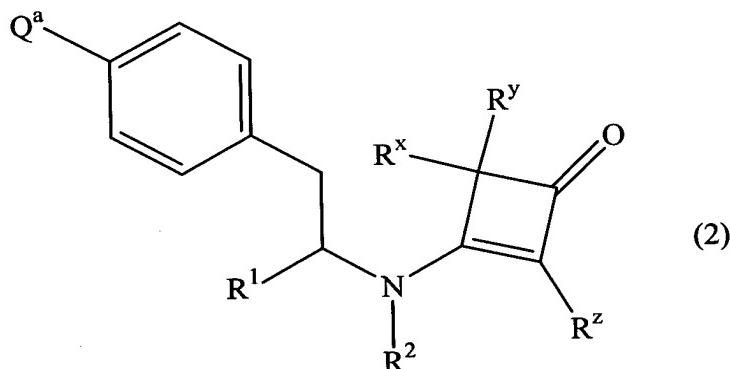
22. (previously presented) A process according to Claim 1 for the preparation of:

ethyl (2S)-2-(2-bromo-3-oxo-spiro[3.5]non-1-en-1-ylamino)-3-[4-([2,7]naphthyridin-1-ylamino)phenyl]propanoate;
and the salts, solvates, hydrates and N-oxides thereof.

23. (previously presented) A process according to Claim 1 for the preparation of:
ethyl (2S)-2-[(2-isopropylsulfanyl-3-oxo-7-oxa-spiro[3.5]non-1-en-1-yl)amino]-3-[4-
([2,7]naphthyridin-1-ylamino)phenyl]propanoate;
and the salts, solvates, hydrates and N-oxides thereof.

24. (previously presented) A process according to Claim 1 for the preparation of:
2-hydroxyethyl (2S)-2-(2-bromo-3-oxo-spiro[3.5]non-1-en-1-ylamino)-3-{4-[(3,5-
dichloroisonicotinoyl)amino]phenyl}propanoate;
and the salts, solvates, hydrates and N-oxides thereof.

25. (currently amended) A compound of formula (2):



wherein:

R¹ is a carboxylic acid (-CO₂H) or a derivative or biostere thereof an acyclic or cyclic carboxylic acid ester, an amide, tetrazole, phosphonic acid, phosphinic acid, sulphonic acid, sulphinic acid, boronic acid, or an acylsulphonamide group;

R² is a hydrogen atom or a C₁₋₆alkyl group;

R^x, R^y and R^z, which may be the same or different, are each an atom or group -L¹(Alk¹)_n(R³)_v in which , or R^z is -L¹(Alk¹)_n(R³)_v and R^x and R^y are joined together to form an optionally substituted spiro linked cycloaliphatic or heterocycloaliphatic group;

L¹ is a covalent bond or a linker atom or group an -O-, -S-, or -Se- atom or an -C(O)-, -C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸).

-N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-,
-N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- group;

R⁸ is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group;

Alk¹ is an optionally substituted aliphatic chain or an optionally substituted heteroaliphatic chain containing one to four -O- or -S- atoms or -C(O)-, -C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸), -N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-, -N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- groups that interrupt or are at the terminus of the aliphatic chain;

R³ is a hydrogen or halogen atom or group selected from -OR^{3a} [where R^{3a} is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group or C₃-cycloalkyl group], -SR^{3a}, -CN and an optionally substituted cycloaliphatic, heterocycloaliphatic, polycycloaliphatic, heteropolycycloaliphatic, aromatic or heteroaromatic group; ;

R^{3a} is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group or C₃-8cycloalkyl group;

n is zero or the integer 1; and

v is the integer 1, 2 or 3;

provided that when n is zero and L¹ is a covalent bond, v is the integer 1;
or R^z is an atom or group as previously defined and R^x and R^y are joined together to form an optionally substituted spiro linked cycloaliphatic or heterocycloaliphatic group;

Q^a is a group -N(R⁴)H;

R⁴ is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group;

and the salts, solvates, hydrates and N-oxides thereof.

26. (original) A compound according to Claim 25 which is:

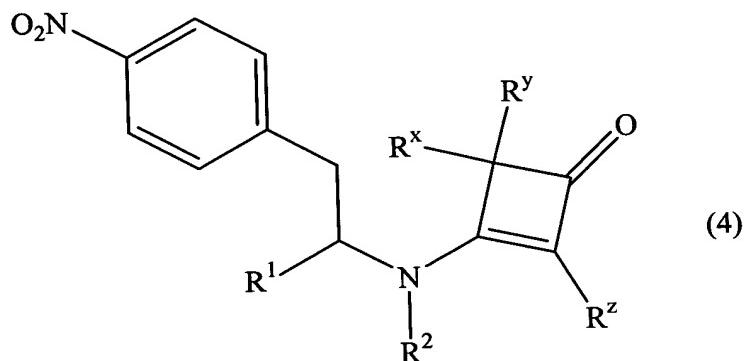
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3-(4-aminophenyl)-2(S)-(3-oxo-7-oxaspiro[3.5]non-1-en-1-ylamino)-propionic acid hydroxyethyl ester.

27. (currently amended) A compound of formula (4):



wherein:

R¹ is a carboxylic acid (-CO₂H) or a derivative or biostere thereof an acyclic or cyclic carboxylic acid ester, an amide, tetrazole, phosphonic acid, phosphinic acid, sulphonic acid, sulphinic acid, boronic acid, or an acylsulphonamide group;

R² is a hydrogen atom or a C₁-6alkyl group;

R^x, R^y and R^z, which may be the same or different, are each an atom or group -L¹(Alk¹)_n(R³)_v in which, or R^z is -L¹(Alk¹)_n(R³)_v and R^x and R^y are joined together to form an optionally substituted spiro linked cycloaliphatic or heterocycloaliphatic group;

L¹ is a covalent bond or a linker atom or group an -O-, -S-, or -Se- atom or an -C(O)-, -C(O)O-, -OC(O)-, -C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸)-, -N(R⁸)CO-, -N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-, -N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- group;

R⁸ is a hydrogen atom or an optionally substituted straight or branched C₁-6alkyl group;

Alk¹ is an optionally substituted aliphatic chain or an optionally substituted heteroaliphatic chain containing one to four -O- or -S- atoms or -C(O)-, -C(O)O-, -OC(O)-,

-C(S)-, -S(O)-, -S(O)₂-, -N(R⁸)-, -CON(R⁸)-, -OC(O)N(R⁸)-, -CSN(R⁸), -N(R⁸)CO-,
-N(R⁸)C(O)O-, -N(R⁸)CS-, -S(O)₂N(R⁸)-, -N(R⁸)S(O)₂-, -N(R⁸)O-, -ON(R⁸)-,
-N(R⁸)N(R⁸)-, -N(R⁸)CON(R⁸)-, -N(R⁸)CSN(R⁸)-, or -N(R⁸)SO₂N(R⁸)- groups that
interrupt or are at the terminus of the aliphatic chain;

R³ is a hydrogen or halogen atom or group selected from -OR^{3a} [where R^{3a} is a hydrogen atom or an optionally substituted straight or branched C₁₋₆alkyl group or C₃₋₈cycloalkyl group], -SR^{3a}, -CN and an optionally substituted cycloaliphatic, heterocycloaliphatic, polycycloaliphatic, heteropolycycloaliphatic, aromatic or heteroaromatic group; ;

R^{3a} is a hydrogen atom or an optionally substituted straight or branched C₁₋₆alkyl group or C₃₋₈cycloalkyl group;

n is zero or the integer 1; and

v is the integer 1, 2 or 3;

provided that when n is zero and L¹ is a covalent bond, v is the integer 1;
~~or R^z is an atom or group as previously defined and R^x and R^y are joined together to form an optionally substituted spiro linked cycloaliphatic or heterocycloaliphatic group;~~
and the salts, solvates, hydrates and N-oxides thereof.

28. (original) A compound according to Claim 27 which is:

3-(4-nitrophenyl)-2(S)-(3-oxo-7-oxaspiro[3.5]non-1-en-1-ylamino)propionic acid hydroxyethyl ester.